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## **CLAIM LISTING:**

Claims 1-59 (Canceled)

60. (Original) A compound of the Formula I

$$R^{1}$$
 $R^{2}$ 
 $R^{3}$ 
 $R^{3}$ 

or a pharmaceutically acceptable salt thereof, wherein:

X<sup>1</sup> is O, NR<sup>4</sup> (where R<sup>4</sup> is hydrogen or alkyl), S, or CR<sup>5</sup>R<sup>6</sup> (where R<sup>5</sup> and R<sup>6</sup> are independently hydrogen or alkyl) or C=O;

Ar1 is aryl or heteroaryl;

 $R^2$  is hydrogen alkyl, acyl, alkoxycarbonyl, aryloxycarbonyl, heteroalkylcarbonyl, heteroalkyloxycarbonyl or  $-R^{21}-R^{22}$  where  $R^{21}$  is alkylene or -C(=0)- and  $R^{22}$  is alkyl or alkoxy;

R<sup>1</sup> is hydrogen, alkyl, haloalkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, cycloalkyl, cycloalkyl, heteroaralkyl, heteroalkylsubstituted cycloalkyl, heterosubstituted cycloalkyl, heteroalkyl, cyanoalkyl, heterocyclylalkyl, R<sup>12</sup>-SO<sub>2</sub>-heterocycloamino (where R<sup>12</sup> is haloalkyl, aryl, aryalkyl, heteroaryl or heteroaralkyl), -Y<sup>1</sup>-C(O)-Y<sup>2</sup>-R<sup>11</sup> (where Y<sup>1</sup> and Y<sup>2</sup> are independently either absent or an alkylene group and R<sup>11</sup> is hydrogen, alkyl, haloalkyl, hydroxy, alkoxy, amino, monoalkylamino or dialkylamino), (heterocyclyl)(cycloalkyl)alkyl or (heterocyclyl)(heteroaryl)alkyl; and

R<sup>3</sup> is hydrogen, alkyl, cycloalkyl, cycloalkylalkyl, aryl, aralkyl, haloalkyl, heteroalkyl, cyanoalkyl, alkylene-C(O)-R<sup>31</sup> (where R<sup>31</sup> is hydrogen, alkyl, hydroxy, alkoxy, amino, monoalkylamino or dialkylamino), amino, monoalkylamino, dialkylamino or NR<sup>32</sup>-Y<sup>3</sup>-R<sup>33</sup> (where Y<sup>3</sup> is -C(O), -C(O)O-, -C(O)NR<sup>34</sup>, S(O)<sub>2</sub> or S(O)<sub>2</sub>NR<sup>35</sup>; R<sup>32</sup>, R<sup>34</sup> and R<sup>35</sup> are independently

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hydrogen or alkyl; and R<sup>33</sup> is hydrogen, alkyl, cycloalkyl, cycloalkylalkyl, heteroalkyl or optionally substituted phenyl) or acyl.

- 61. (Original) The compound of Claim 1, wherein Ar<sup>1</sup> is optionally substituted phenyl.
  - 62. (Original) The compound of Claim 61, wherein X<sup>1</sup> is O or CH<sub>2</sub>.
  - 63. (Original) The compound of Claim 62, wherein  $X^1$  is O.
- 64. (Original) The compound of Claim 63 wherein R<sup>1</sup> is aryl, aralkyl, cycloalkyl, cycloalkyl, heteroalkylsubstituted cycloalkyl, heterosubstituted cycloalkyl, heterocyclyl or heterocyclyl alkyl.
- 65. (Original) The compound of Claim 64, wherein R<sup>1</sup> is heteroalkylsubstituted cycloalkyl, heteroalkyl, heteroalkyl, heteroalkyl or heterocyclyl.
  - 66. (Original) The compound of Claim 65, wherein R<sup>1</sup> is heterocyclyl.
  - 67. (Original) The compound of Claim 65, wherein R<sup>1</sup> is heteroalkyl.
  - 68. (Original) The compound of Claim 67, wherein R<sup>1</sup> is hydroxyalkyl.
- 69. (Original) The compound of Claim 65, wherein Ar<sup>1</sup> is 2-substituted-phenyl, 4-substituted-phenyl or 2,4-disubstituted-phenyl.
- 70. (Original) The compound of Claim 69, wherein Ar<sup>1</sup> is 2-chlorophenyl, 2-fluorophenyl, 2-fluoro-4-methylphenyl, 4-fluoro-2-methyl or 2,4-difluorophenyl.
  - 71. (Original) The compound of Claim 70, wherein R<sup>3</sup> is methyl.

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- 72. (Original) The compound of Claim 71, wherein R<sup>1</sup> is heteroalkyl substituted cycloalkyl, heteroalkyl, heteroalkyl or heterocyclyl.
- 73. (Original) The compound of Claim 72, wherein R<sup>1</sup> is heterocyclyl.
- 74. (Original) The compound of Claim 72, wherein R<sup>1</sup> is heteroalkyl.
- 75. (Original) The compound of Claim 72, wherein R<sup>1</sup> is hydroxyalkyl.
- 76. Currently Amended) A method for treating p38 mediated disorder arthritis, said method comprising administering to a patient in need of such treatment, an effective amount of a compound of Claim 60.
  - 77. (Canceled)

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